

**510(k) Summary  
for the Gridlock Plating System**

**MAY 08 2013**

In accordance with 21 CFR 807.92 of the Federal Code of Regulations the following 510(k) summary is submitted for  
the Gridlock Plating System

**1. GENERAL INFORMATION**

**Date Prepared:** April 3, 2013

**Trade Name:** Gridlock Plating System

**Common Name:** bone plate & screws

**Classification Name:** Single/multiple component metallic bone fixation appliances and accessories  
Smooth or threaded metallic bone fixation fastener

**Class:** II

**Product Code:** HRS / HWC

**CFR section:** 21 CFR section 888.3030 / 888.3040

**Device panel:** Orthopedic

**Legally Marketed:** Gridlock Plating System (K121452, K123525)

**Predicate Device:** OsteoMed Foot Plating System (K091614)

**Submitter:** Trilliant Surgical LTD  
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Houston, TX 77024  
1-800-495-2919 Tele

**Contact:** J.D. Webb  
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Round Rock, TX 78681  
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**2. DEVICE DESCRIPTION**

Gridlock Plating System consists of various shape and size plates for the management of orthopedic osteotomies, reconstruction, and trauma. Features include a low profile, limited contact, capability of dynamic/manual compression, and angulated-locking threaded screw holes. The system also consists of multiple locking/standard screw lengths and diameters and the necessary instruments to facilitate the placement of these implants.

**Change from Predicate:**

This Special 510(k) is submitted in order to gain clearance for additional plates and screws in the Gridlock Plating System.

**Materials:**

CP Titanium per ASTM F67  
Titanium alloy per ASTM F136

**3. INTENDED USE**

The Gridlock Plating System is intended for use in trauma and reconstructive procedures of the small bones in the hand/foot, ankle, and other bones appropriate for the size of the device.

The plates (implant), screws (implant), olive wires (instrument), and guide wires (instrument) are intended for single use only.

**4. SUMMARY OF TECHNOLOGICAL CHARACTERISTICS / SUBSTANTIAL EQUIVALENCE**

The Gridlock Plating System is substantially equivalent to the predicate devices in terms of intended use, design, materials used, mechanical safety and performances. Two bone plates have been added to the Gridlock Plating System. These two plates are for use in treating the first metatarsal phalangeal joint and are manufactured from the same material, use the same screws and instruments, and are packaged and sterilized the using the same methods as the predicate Gridlock plates.

**5. NON-CLINICAL TEST SUMMARY**

Engineering analysis was performed.

**6. CLINICAL TEST SUMMARY**

No clinical studies were performed

**7. CONCLUSIONS NONCLINICAL AND CLINICAL**

Trilliant Surgical LTD considers the current Gridlock Plating System to be equivalent to the predicate devices listed above. This conclusion is based upon the devices' similarities in principles of operation, technology, materials and indications for use.



Food and Drug Administration  
10903 New Hampshire Avenue  
Document Control Center – WO66-G609  
Silver Spring, MD 20993-0002

May 8, 2013

Trilliant Surgical LTD  
% The OrthoMedix Group, Incorporated  
Mr. J.D. Webb  
Authorized Correspondent  
1001 Oakwood Boulevard  
Round Rock, Texas 78681

Re: K130964

Trade/Device Name: Gridlock Plating System

Regulation Number: 21 CFR 888.3030

Regulation Name: Single/multiple component metallic bone fixation appliances and accessories

Regulatory Class: Class II

Product Code: HRS, HWC

Dated: April 3, 2013

Received: April 8, 2013

Dear Mr. Webb:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA).

You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21

CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address <http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to <http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm> for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address <http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>.

Sincerely yours,

**Erin D. Keith**

Mark N. Melkerson  
Director  
Division of Orthopedic Devices  
Office of Device Evaluation  
Center for Devices and  
Radiological Health

Enclosure

## INDICATIONS FOR USE

510(k) Number (if known): K130964

Device Name: Gridlock Plating System

### Indications for Use:

The Gridlock Plating System is intended for use in trauma and reconstructive procedures of the small bones in the hand/foot, ankle, and other bones appropriate for the size of the device.

The plates (implant), screws (implant), olive wires (instrument), and guide wires (instrument) are intended for single use only.

Prescription Use X  
(Part 21 CFR 801 Subpart D)

AND/OR

Over-The-Counter Use \_\_\_\_\_  
(21 CFR 807 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE OF NEEDED)

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Concurrence of CDRH, Office of Device Evaluation (ODE)

**Elizabeth M. Frank -S**

Division of Orthopedic Devices